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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/016,964	12/14/2001	Heidi Riedel	Beiersdorf 755-KGB	7321	
7055 GREENBLUM	7590 07/02/2010 I & BERNSTEIN, P.L.C		EXAMINER		
1950 ROLANI	D CLARKE PLACE	•	KANTAMNENI, SHOBHA		
RESTON, VA	20191		ART UNIT	PAPER NUMBER	
			1627		
			NOTIFICATION DATE	DELIVERY MODE	
			07/02/2010	ELECTRONIC	

## Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

gbpatent@gbpatent.com pto@gbpatent.com

## Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)		
10/016,964	RIEDEL ET AL.		
Examiner	Art Unit		
Shobha Kantamneni	1627		
Onobila Namamilem	1027		

	Shobha Kantamneni	1627	
The MAILING DATE of this communication appe	ars on the cover sheet with the	correspondence add	ress
THE REPLY FILED 14 June 2010 FAILS TO PLACE THIS APP	LICATION IN CONDITION FOR A	LLOWANCE.	
<ol> <li>X The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Appe for Continued Examination (RCE) in compliance with 37 C periods:</li> </ol>	replies: (1) an amendment, affidavi eal (with appeal fee) in compliance	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request
a) The period for reply expiresmonths from the mailing b) The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire Is Examiner Note: If box 1 is checked, check either box (a) or	dvisory Action, or (2) the date set forth ater than SIX MONTHS from the mailing	date of the final rejection	n.
MONTHS OF THE FINAL REJECTION. See MPEP 706.07(	).		
Extensions of time may be obtained under 37 CFR 1.136(a). The date have been filed is the date for purposes of determining the period of ext under 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked: Any reply neceived by the Office later may reduce any earned patient term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	on which the petition under 37 CFR 1.1 ension and the corresponding amount hortened statutory period for reply origi than three months after the mailing dat	of the fee. The appropria nally set in the final Offic e of the final rejection, en	ate extension fee e action; or (2) as en if timely filed,
<ol> <li>The Notice of Appeal was filed on A brief in comp filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed w</li> </ol>	sion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	
<u>AMENDMENTS</u>			
<ol> <li>The proposed amendment(s) filed after a final rejection, t         <ul> <li>(a) They raise new issues that would require further cor</li> <li>(b) They raise the issue of new matter (see NOTE belo</li> </ul> </li> </ol>	nsideration and/or search (see NO) w);	TE below);	
(c) They are not deemed to place the application in bet	ter form for appeal by materially red	ducing or simplifying th	ne issues for
appeal; and/or  (d) ☐ They present additional claims without canceling a c	corresponding number of finally reje	ected claims.	
NOTE: (See 37 CFR 1.116 and 41.33(a)).			
4. The amendments are not in compliance with 37 CFR 1.12		mpliant Amendment (F	PTOL-324).
5. Applicant's reply has overcome the following rejection(s):			
<ol> <li>Newly proposed or amended claim(s) would be all non-allowable claim(s).</li> </ol>	owable if submitted in a separate, i	imely filed amendmer	it canceling the
<ol> <li>For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is provided.</li> </ol>		I be entered and an ex	xplanation of
The status of the claim(s) is (or will be) as follows: Claim(s) allowed: NONE.			
Claim(s) objected to:			
Claim(s) rejected: <u>18-41,44 and 45</u> . Claim(s) withdrawn from consideration:			
AFFIDAVIT OR OTHER EVIDENCE			
<ol> <li>The affidavit or other evidence filed after a final action, bu because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e).</li> </ol>			
<ol> <li>The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to o showing a good and sufficient reasons why it is necessary</li> </ol>	vercome <u>all</u> rejections under appea	al and/or appellant fails	to provide a
10. The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER	n of the status of the claims after er	ntry is below or attache	ed.
<ol> <li>The request for reconsideration has been considered bu See page 2.</li> </ol>	does NOT place the application in	condition for allowan	ce because:
12. Note the attached Information Disclosure Statement(s). ( 13. Other:	PTO/SB/08) Paper No(s)		
	/Shengjun Wang/ Primary Examiner, Art U	nit 1627	

Continuation of 11: Applicant's arguments have been considered, but not found persuasive as discussed in the previous office actions, and those found below. All the rejections made in the final office action are maintainted.

The rejection of Claims 18-20, 25, 27-33, 35, 36 under 35 U.S.C. 102(b) as being anticipated by Beutler et al. (US 4,808,388, PTO-1449).

Applicant argues that "the "Polysorbate 20" employed in Examples 5/la and 5/lb of BEUTLER is expressly identified in BEUTLER as "Sorbitan monolaurate" (see Example 5/la), i.e., not as polyocythylene(20) sorbitan monolaurate as alleged by the Examiner. As can be taken from, e.g., http://www.fac.org/lag/agn/iecfa- additives/specs/Monographl/Additive-431.pdf, sorbitan monolaurate is a commercial product which finds use as emulsifier." These arguments have been considered, but not found persuasive been Beutler clearly discloses in Example 5/1b, the employment of polysorbate 20 as the nonionic emulsifier. Polysorbate 20 is a polyoxyethylene derivative of sorbitan monolaurate, and not just sorbitan monolaurate. See

http://chemicalland21.com/specialtychem/perchem/POLYOXYETHYLENE%20SORBITAN%20LAURATE.htm

Applicant argues that "It is not seen that the preparations of Examples 5/Ia and 5/Ib of BEUTLER necessarily comprise from 1% to 90 % by volume of at least one gas".

These remarks have been considered. It is pointed out that the amount of the gaseous propellant is about 1 to 4 % by weight. See column 4. lines 34-44.

Beutler et al. in Example 5/la and Example 5/lb discloses a composition comprising 5 % by weight of TEA-Stearate (Triethanolamine Stearate) i.e. wholloy or partially neutralized fatty acid instant emulsifier A. 0, 5 % by weight of 19EG 20 sorbitan monolaurate) i.e. instant emulsifier B. 1% by weight of cetyl alcohol i.e. instant emulsifier B, 1% by weight of oretyl alcohol i.e. instant emulsifier B, 1% by weight of 10 to 4% by weight of 20 kept alcohol i.e. instant emulsifier B, and in an amount of 1 to 4 % by weight, which meets the instant claims.

Applicant's remarks made regarding NetMoneyIN, Inc. v. VeriSign, Inc., 545 F.3d 1359 (Fed. Cir. 2008), it is pointed out that as discussed above Beutler et al. discloses all the elements arranged as in instant composition as in instant Claim 18

The rejection of claims 18-24, 28-31, 34, 36-39, 44 under 35 U.S.C. 103(a) as being unpatentable over Bellon et al. (FR 2,789,397 with English translation of record):

Applicant's arguments regarding PEG-100 stearate gloceryl stearate in Example 1 of BELLON (marketed by SEPPIC Company as glyceryl stearate from SEPPIC contains an ester of stearia and PEG-100 stearate) has been considered, it is pointed out again PEG-100 stearate/glyceryl stearate from SEPPIC contains an ester of stearia acid and polyethylene glycol, PEG-100 refers to polyethylene glycol comprising 100 ethylene glycol units. Thus, PEG-100 stearate glyceryl stearate taught by BELLON in Example 1 reads on instant emulsifier B which is a polyethylene glycol units. Thus, PEG-100 stearate glyceryl stearate taught by BELLON in Example 1 reads on instant emulsifier B which is a polyethylene glycol units. Thus, PEG-100 stearate glyceryl stearate http://www.gapregnancy.com/sourcest/gu/2009/montanov-68.pdf

Applicant argues that "BELLON does not even discuss the combination of compounds which allegedly correspond to the present enuisifiers A, B and C, let alone as an emulsifier system, but mentions these compounds separately and for unrelated purposes or not at all." These arguments have been considered, but not found persuasive. Bellon et al. exemplify a facial foam composition or preparation comprising 22% PEG-100 stearate/glyceny is bearate combination from SEPPIC which is a polyethoxylatef acid ester in the instant claim 18, componant B, stearate having a chain 18 carbons and 100 of ethoxylation, 12% stearic acid which is a fatty adol in the instant claim 18, component A, stearic acid having a chain 18 carbons; 6% octylodocanol, which is a fatty alcohol in the instant claim 18, component C having a chain 20 carbons; nitrogen added to the composition in 70% by volume which is one gas in claim 18, and thus meets instant claim limitation 18. See Example 1 and Table 1 (at pase 10-11) and 16 of the English translation.

Applicant argues that "the Examiner has failed to provide any evidence that dept glucoside, i.e., and ther of glucose and decanol, is substantially the same as a failty alcohol (such as, e.g., the oxyldodecanol employed in Example 1 of BELCON). These arguments have been considered, but not found persuasive. It is pointed out that Example 1 of Bellon comprises oxyldodecanol i.e a fatty alcohol, and thus meets instant dain 18 limitation.

The rejection of claims 18-20, 24-25, 28, 30-31, 33-36, 44 are rejected under 35 U.S.C. 102(b) as being anticipated by Penska et al. (EP 0 938 890 or US 5,851,544, PTO-1449, IDS filed on 06/20/2006).

Appleant argument regarding self-foaming or foam-like have been considered, but not found persuasive because Penska discloses instant composition as in claim 18, and thus meets the instant claim limitation.

Applicant argues that "Applicants do not know how the Examiner has calculated the alleged volume percentage of perfluorodecane in the composition of Example 6 of PENSKA based on the indicated weight percentage thereof". It is pointed out that Example 6, 50 wt % of perfluorodecane corresponds to 22.5 % by volume of perfluorodecane, since the density of perfluorodecane = 1.7.